

U.S. Label Action Checklist and Approval Form

To initiate this form, do the following (DO NOT delete or alter any section of this form):

- Place an X in the box under Type of Action (see below).
- Fill in product name, active ingredient(s), and EPA registration number.
- If form is for a Section 18 or 24(c) label, fill out the appropriate section. If form is not for Section 18 or 24(c) label, ignore these sections.
- Fill in the person's name next to the appropriate role for the product.
- List the label changes on the last page of the form.
- Send the form via e-mail to the Regulatory Specialist for the product.

To approve this form, do the following:

- Fill out the Key Questions/Information Required for your role.
- Put the date of your approval in the Approval Date column on the same line as your name.

Note: This form is to approve the proposed action, not review/approve the draft or proposed label. All sections of this form must be completed prior to initiation of draft labeling by the Regulatory Specialist.

Type of Action: (place an X in the box to the left of the appropriate action)

[Creation of Section 3 main label	[Creation of Section 3 supplemental label
F		FO	
O		RM	
R		CH	
M		EC	
C		KB	
H		OX	
E]	
C			
K			
B			
O			
X			
]			
[Creation of Section 18 label	[Creation of Section 24(c) label
F		FO	
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R		CH	
M		EC	
C		KB	
H		OX	
E]	
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[Creation of Section 2(ee) label		
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Amendment of Section 3 main label

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Amendment of Section 3 supplemental label

Amendment of Section 18 label

[
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EC
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Amendment of Section 24(c) label

Amendment of Section 2(ee) label

Product Name: [PAGE][FORMTEXT] a.i.: [FORMTEXT]
EPA Reg. No.: [FORMTEXT]

If Section 18, what is the objective: [FORMTEXT]

If Section 18, who is sponsoring (e.g., state, commodity groups): [FORMTEXT]

If Section 24(c), what is the objective: [FORMTEXT]

Approval: (include date in Approval Date column in the row containing your name)

Role	Name	Input/Comments	Approval Date
Product Technology Specialist	[FORMTEXT]	[FORMTEXT]	[FORMTEXT]
Marketing Specialist	[FORMTEXT]	[FORMTEXT]	[FORMTEXT]

Tech Expert	[FORMTEXT]	[FORMTEXT]	[FORMTEXT]
Federal Regulatory Manager	[FORMTEXT]	[FORMTEXT]	[FORMTEXT]
State Regulatory Manager	[FORMTEXT]	[FORMTEXT]	[FORMTEXT]

Key Questions/Information Required (Information Provided by Tiger Team):

Product Technology Specialist

• Use directions for proposed use or list of proposed label changes, including those listed on label bulletin board (Yes/No): (Provide text/create list on page 2 or embed Word document on page 2)	[FORMTEXT]
• Sec 18/24(c) labels only: Include Special Conditions of Use (which was previously called Waiver of Liability) on label (Yes/No):	[FORMTEXT]

Marketing Specialist:

• Reason for proposed change(s) or creation of label:	[FORMTEXT]
• ♦ Mandatory or Regulatory initiated (e.g., PR Notice, RED, state req., etc.):	[FORMTEXT]
• ♦ Commercial (new crop, rates, pest, etc.): (New crop, rates, pest will be submitted as a supplemental label)	[FORMTEXT]
• Molecule status per current year Business Line Matrix (Invest, Grow, Defend, Manage for Cash):	[FORMTEXT]
• Projected market value of proposed change(s) or use(s):	\$(FORMTEXT]
• List of state(s) to be registered in:	[FORMTEXT]
• Expected/desired sales promotion date:	Date: [FORMTEXT]
• Sec 3 main label only: Does action involve a supplemental distributor (Yes/No):	[FORMTEXT]
• If yes, will DAS commercialize label too (Yes/No):	[FORMTEXT]
• Sec 3 main label only: Will there be any post-approval label activities, e.g. registration of ABN, roll same changes onto ABN label, a subset of registered uses commercialized (Federal Regulatory Manager to be consulted) (Yes/No)	[FORMTEXT]
• If yes, provide specific details:	[FORMTEXT]

Technical Expert (Review and Comment with Input from Product Technology Specialist):

• Availability of efficacy and/or crop safety data required by Cal DPR?: (Data must be supplied to Regulatory (State Regulatory Assistant) before label before product(s) can be registered in the states.)	[FORMTEXT]
• Has risk/benefit review been conducted (e.g., crop safety, efficacy) (Yes/No):	[FORMTEXT]
• Risk score from Decision Tree Risk Matrix (≤ 45 , between 45 and 65, >65): (If score is greater than 65, Special Conditions of Use language may be required.)	[FORMTEXT]

Federal Regulatory Manager:

• If changes are mandatory, can other changes be made simultaneously (Yes/No):	[FORMTEXT]
• Any outstanding data requirements at EPA (Yes/No):	[FORMTEXT]
• Sec 3 labels only: Last EPA label submission date: (Note: No overlapping label amendments are allowed.)	Date: [FORMTEXT]
• Sec 3 labels only: Target EPA submission date: (Note: Nominal review time is 90 days for amendments requiring no data; up to 2 years for new crops with PRIA.)	Date: [FORMTEXT]
• Is proposed use consistent with federal regulatory strategy (Yes/No):	[FORMTEXT]
• Amendment cost – EPA fee for service:	\$(FORMTEXT]
• Sec 3 main label only: Will there be any post-approval label activities, i.e.,	[FORMTEXT]

registration of ABN, roll same changes onto ABN label, a subset of registered uses commercialized (Marketing Specialist to be consulted) (Yes/No)	
If yes, provide specific details:	[FORMTEXT]

State Regulatory Manager:

• List any special considerations (e.g., PPE for urban pest, special state agreements, etc.):	[FORMTEXT]
• State registration fees associated with this action:	\$(FORMTEXT)
• Any outstanding data requirements in the state(s):	[FORMTEXT]
• All necessary data/information will be assembled and available to support application: (Data must be supplied to Regulatory (State Regulatory Assistant) before product(s) can be registered in the states.)	[FORMTEXT]
• Sec 18/24(c)/2(ee) labels only: Target submission date to state	Date: [FORMTEXT]
• Sec 18 label only: Is there pending Section 3 (Yes/No):	[FORMTEXT]
♦ If so, anticipated approval date:	Date: [FORMTEXT]

Listing of proposed label change concepts to be provided by Product Technology Specialist (e.g., addition of a pest, change of buffer zone, application timing etc.)

1. [FORMTEXT]
 2. [FORMTEXT]
 3. [FORMTEXT]
- [FORMTEXT]